

*a<sub>1</sub>* cont'd

3                   taking a plurality of different crystallization samples in an enclosed microvolume,  
4                   the plurality of crystallization samples comprising a material to be crystallized and crystallization  
5                   conditions which vary among the plurality of crystallization samples;

6                   allowing crystals of the material to form in the plurality of crystallization samples;  
7                   and

8                   identifying which of the plurality of crystallization samples comprise a precipitate  
9                   or a crystal of the material.

*a<sub>2</sub>*

1                 6. (Amended) A method according to claim 1 wherein the enclosed microvolume is  
2                 at least partially defined by a face of a card shaped substrate.

*a<sub>3</sub>*

1                 9<sub>14</sub>. (Amended) A method according to claim 1, the method further comprising  
2                 performing a spectroscopic analysis on a precipitate or crystal formed within the microvolume.

10

1                 15. (Amended) A method according to claim 14, wherein the spectroscopic analysis  
2                 is selected from the group consisting of Raman, UV/VIS, IR, and x-ray spectroscopy.

*a<sub>4</sub>*

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1                 17. (Amended) A method according to claim 16, wherein x-ray spectroscopy is  
2                 performed such that a portion of the microvolume that the x-ray beam traverses contains at least  
3                 as many electrons as is contained in a material defining the portion of the microvolume that the  
4                 x-ray beam traverses.

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1                 18. (Amended) A method according to claim 16, wherein x-ray spectroscopy is  
2                 performed such that a portion of the microvolume that the x-ray beam traverses contains at least  
3                 three times as many electrons as is contained in a material defining the portion of the  
4                 microvolume that the x-ray beam traverses.

*a<sub>5</sub>*

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1                 19. (Amended) A method according to claim 16, wherein x-ray spectroscopy is  
2                 performed such that a portion of the microvolume that the x-ray beam traverses contains at least

*A<sub>3</sub> cont'd*

3 five times as many electrons as is contained in a material defining the portion of the  
4 microvolume that the x-ray beam traverses.

*15*  
1 20. (Amended) A method according to claim 16, wherein x-ray spectroscopy is  
2 performed such that a portion of the microvolume that the x-ray beam traverses contains at least  
3 ten times as many electrons as is contained in a material defining the portion of the microvolume  
4 that the x-ray beam traverses.

*16*  
1 21. (Amended) A method according to claim 1, wherein material defining the  
2 microvolume defines a groove that reduces a number of electrons that an x-ray beam used to  
3 perform x-ray spectroscopy of a crystal within the microvolume traverses in the process of  
4 performing x-ray spectroscopy on the sample within the microvolume.

*A<sub>5</sub>*  
1 19 24. (Amended) A method according to claim 1, wherein one or more dividers are  
2 positioned within the enclosed microvolume to separate adjacent crystallization samples within  
3 the enclosed microvolume.

*Sub 23*  
1 25. (Amended) A method according to claim 25, wherein the  
2 one or more dividers are formed of an impermeable material.

*Sub 24*  
1 28. (Amended) A method according to claim 25, wherein the  
2 one or more dividers are formed of a permeable material.

1 29. (Amended) A method according to claim 25, wherein the  
2 one or more dividers are formed of a semipermeable material.

*A<sub>7</sub>*  
1 33. (Amended) A method according to claim 25, wherein at least one of the one or  
2 more dividers form an interface selected from the group consisting of liquid/liquid, liquid/ gas  
3 interface, liquid/ solid and liquid/ sol-gel interface.

*cont'd*

1        34. (Amended) A method according to claim 25, wherein the one or more dividers  
2 are selected from the group consisting of a membrane, gel, frit, and matrix.

1        35. (Amended) A method according to claim 25, wherein the one or more dividers  
2 function to modulate diffusion characteristics between adjacent crystallization samples.

1        36. (Amended) A method according to claim 25, wherein at least one of the one or  
2 more dividers is formed of a semipermeable material which allows diffusion between adjacent  
3 crystallization samples.

1        37. (Amended) A method for determining crystallization conditions for a material,  
2 the method comprising:

3                taking a plurality of different crystallization samples in a plurality of enclosed  
4 microvolumes, each microvolume comprising one or more crystallization samples, the  
5 crystallization samples comprising a material to be crystallized and crystallization conditions  
6 which vary among the plurality of crystallization samples;

7                allowing crystals of the material to form in the plurality of crystallization samples;  
8 and

9                identifying which of the plurality of crystallization samples comprise a precipitate  
10 or a crystal of the material.

Please add the following new claims 38-45.

*38.* A method according to claim *16*, wherein the x-ray spectroscopy is x-ray  
2 diffraction.

*39.* A method according to claim *16*, wherein x-ray spectroscopy is performed such  
2 that a portion of the crystal or precipitate that the x-ray beam traverses contains at least as many  
3 electrons as is otherwise traversed by the x-ray beam when traversing a device comprising the  
4 microvolume.

*34*

1 40. A method according to claim 16, wherein x-ray spectroscopy is performed such  
2 that a portion of the crystal or precipitate that the x-ray beam traverses contains at least three  
3 times as many electrons as is otherwise traversed by the x-ray beam when traversing a device  
4 comprising the microvolume.

*35*

1 41. A method according to claim 16, wherein x-ray spectroscopy is performed such  
2 that a portion of the crystal or precipitate that the x-ray beam traverses contains at least five times  
3 as many electrons as is otherwise traversed by the x-ray beam when traversing a device  
4 comprising the microvolume.

*36*

1 42. A method according to claim 16, wherein x-ray spectroscopy is performed such  
2 that a portion of the crystal or precipitate that the x-ray beam traverses contains at least ten times  
3 as many electrons as is otherwise traversed by the x-ray beam when traversing a device  
4 comprising the microvolume.

*37*

1 43. A method according to claim 37, wherein each microvolume comprising a  
2 plurality of crystallization samples.

*38*

1 44. A method according to claim 16, wherein x-ray spectroscopy is performed such  
2 that a portion of the microvolume that the x-ray beam traverses contains at least half as many  
3 electrons as is contained in a material defining the portion of the microvolume that the x-ray  
4 beam traverses.

*39*

1 45. The method according to claim 1 wherein the material to be crystallized contains  
2 at least two or more materials selected from the group consisting of viruses, proteins, peptides,  
3 nucleosides, nucleotides, ribonucleic acids, deoxyribonucleic acids, small molecules, drugs,  
4 putative drugs, inorganic compounds, metal salts, organometallic compounds and elements.